<u>Listing of Claims</u>:

- 177. The therapeutic composition of claim 210 or 211, wherein said genetically-engineered monoclonal antibody is a single-chain antibody.
 - 210. A therapeutic composition, comprising: a pharmaceutical formulation comprising
 - (1) a pharmaceutically acceptable carrier and
- (2)(a) a genetically-engineered antibody that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid, or
- (b) a fragment of the genetically-engineered antibody of (a) that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid,

wherein said genetically-engineered antibody is obtained by genetically engineering the DNA encoding a monoclonal antibody that

- (i) inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid and
- (ii) is obtainable using a peptide consisting of residues 1-28 of beta-amyloid as an immunogen or recognizes an epitope within residues 1-28 of beta-amyloid.
- 211. The therapeutic composition of claim 210, wherein said genetically-engineered antibody of (2)(a) inhibits aggregation of human beta-amyloid or maintains the solubility of soluble human beta-amyloid, or said fragment of (2)(b) inhibits aggregation of human beta-amyloid or maintains the solubility of soluble human beta-amyloid, and said

genetically-engineered antibody of (2)(a) is obtained by genetically engineering the DNA encoding a monoclonal antibody that inhibits aggregation of human beta-amyloid or maintains the solubility of soluble human beta-amyloid and said monoclonal antibody is obtainable using a peptide consisting of residues 1-28 of human beta-amyloid as an immunogen or recognizes an epitope within residues 1-28 of human beta-amyloid.

- 212. A therapeutic composition, comprising: a pharmaceutical formulation comprising
- (1) a pharmaceutically acceptable carrier and
- (2)(a) a human monoclonal antibody that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid, or
- (b) a fragment of the human monoclonal antibody of (a) that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid,

wherein said human monoclonal antibody is obtainable using a peptide consisting of residues 1-28 of beta-amyloid as an immunogen.

213. The therapeutic composition of claim 212, wherein said human monoclonal antibody of (2)(a) inhibits aggregation of human beta-amyloid or maintains the solubility of soluble human beta-amyloid, or said fragment of (2)(b) inhibits aggregation of human beta-amyloid or maintains the solubility of soluble human beta-amyloid, and wherein said human monoclonal antibody of (a) is obtainable using a peptide

consisting of residues 1-28 of human beta-amyloid as an immunogen.

214. A method of making a therapeutic composition comprising (1) a pharmaceutically acceptable carrier and (2)(a) a genetically-engineered antibody that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid, or (b) a fragment of the genetically-engineered antibody of (a), which fragment inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid, said method comprising:

selecting a monoclonal antibody that

- (i) inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid, and
- (ii) is obtainable using a peptide consisting of residues 1-28 of beta-amyloid as an immunogen or recognizes an epitope within residues 1-28 of beta-amyloid;

genetically engineering the DNA encoding said selected monoclonal antibody so as to produce a genetically-engineered antibody that inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid, or a fragment of a genetically engineered antibody, which fragment inhibits aggregation of beta-amyloid or maintains the solubility of soluble beta-amyloid; and

formulating said genetically engineered monoclonal antibody or fragment with a pharmaceutical carrier into a pharmaceutical formulation that is a therapeutic composition.